



California Medical Device Recall Information



Recall Name

Verathon Inc. Recalls GlideScope Video Laryngoscope Reusable Blades Due to Breaking Blades

Recall Date	Product Description	Recalling Firm	Recall Reason
5/10/13	Reusable Blades for Video Laryngoscope systems <ul style="list-style-type: none">GlideScope GVLGlideScope AVL	Verathon, Inc. Bothell, WA	<i>Prone to developing cracks and/or breaking off across the tip of the blade.</i>
Recall Class	Product Identification	Distribution	Affected Dates
I	<ul style="list-style-type: none">GlideScope GVL 3 Part # 0574-0007 S/Ns: MD112388 to MD121908GlideScope GVL 4 Part #0574-0001 S/Ns: LG112759 to LG122582GlideScope GVL 5 Model #0574-0030 S/Ns: XL111799 to XL121759GlideScope AVL 2 Part # 0574-0118 S/Ns: AD111500 to AD121604GlideScope AVL 3 Part # 0574-0115 S/Ns: AD111500 to AD121688GlideScope AVL 4 Part # 0574-0116 S/Ns: AE111500 to AE121778GlideScope AVL 5 Part # 0574-0117 S/Ns: AF111500 to AF121666	CA , worldwide	Manufactured and distributed from August 1, 2011 to June 30, 2012. The reusable blades can be identified by the serial number engraved on the label found on the handle of the blade.

FOR ADDITIONAL INFORMATION, PLEASE VISIT:

<http://www.fda.gov/MedicalDevices/Safety/ListofRecalls/ucm360249.htm>